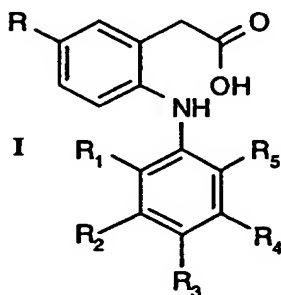


CLAIMS

1. process for the production of a compound of Formula I, or a pharmaceutically acceptable salt thereof, or a pharmaceutically acceptable prodrug ester thereof,



wherein R is methyl or ethyl;

R₁ is chloro or fluoro;

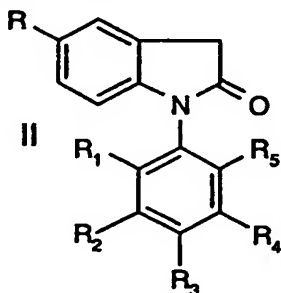
R₂ is hydrogen or fluoro;

R₃ is hydrogen, fluoro, chloro, methyl, ethyl, methoxy, ethoxy or hydroxy;

R₄ is hydrogen or fluoro; and

R₅ is chloro, fluoro, trifluoromethyl or methyl,

provided that R₁, R₂, R₄ and R₅ are not all fluoro when R is ethyl and R₃ is H; comprising cleaving a lactam of formula II

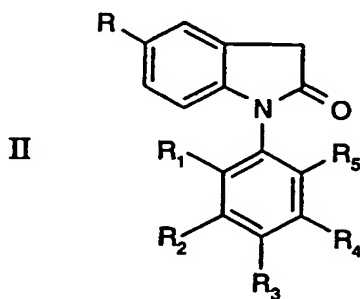


wherein the symbols are as defined above with a base; and in the above process, if desired, temporarily protecting any interfering reactive groups and then isolating the resulting compound of the invention; and, if desired, converting the free carboxylic acid of the compound of formula I into a pharmaceutically acceptable ester derivative thereof; and/or if desired,

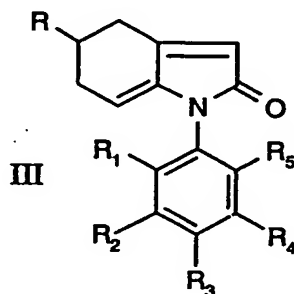
converting the free acid of formula I into a salt or a resulting salt into the free acid or into another salt.

2. A process selected from

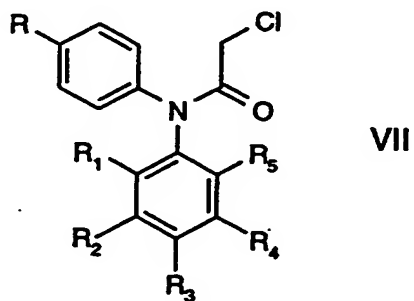
a) a process for the production of a lactam of formula II



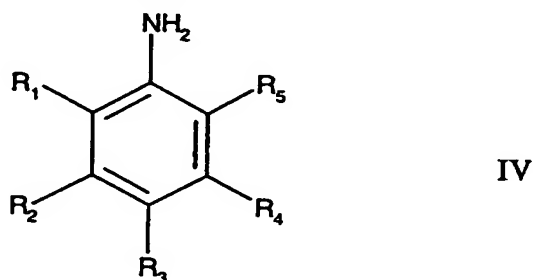
which comprises oxidizing of a lactam of formula III



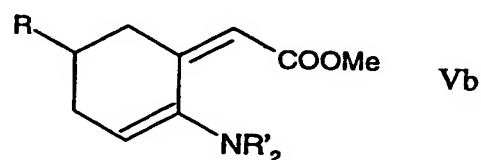
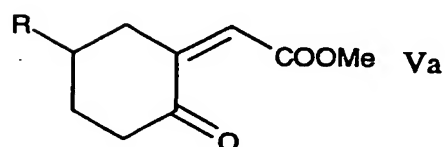
b) a process for the production of the lactam of formula II as defined in a) above, which comprises cyclisation of a compound of formula VII



- c) a process for the preparation of a compound of formula III as defined above comprising coupling as aniline derivative of formula IV

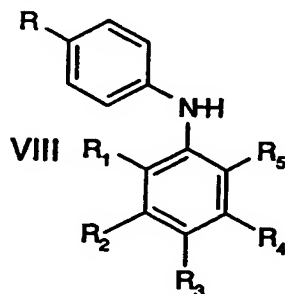


with a cyclohexanone derivative of formula Va or an amino substituted cyclohexene derivative of formula Vb



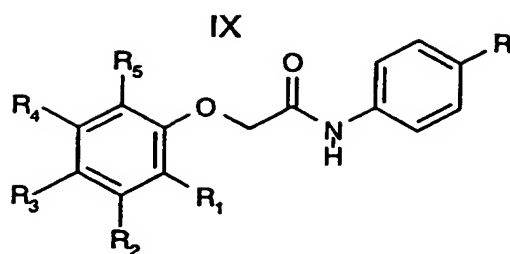
wherein R is ethyl or methyl and R' is lower alkyl or similar or NR'₂ forms a ring as in piperidine or morpholine.

- d) a process for the production of a compound of formula VII which comprises N-acylation of a diphenylamine of formula VIII

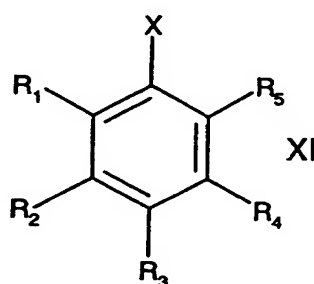


with a haloacetyl chloride

- e) a process for the preparation of a compound of formula VIII which comprises rearrangement and hydrolysis of a compound of formula IX.

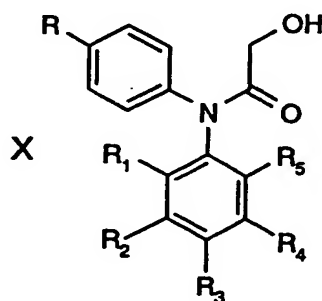


- f) a process for the production of a compound of formula VIII which comprises coupling of a halobenzene derivative of formula XI with p-toluidine or 4-ethyl-aniline

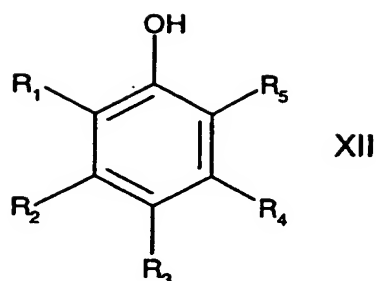


where X is a halogen

- g) a process for the production of a compound of formula VIII as defined in d) above which comprises coupling an aniline derivative of formula IV as defined in c) above with 4-bromotoluene or 1-ethyl-4-bromobenzene
- h) a process for the production of a compound of formula VIII as defined in d) above which comprises cleavage of a compound of formula X

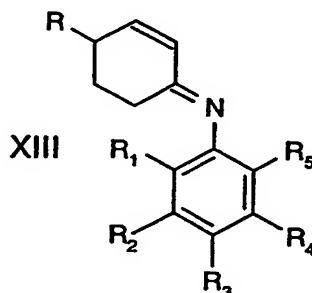


- i) a process for the formation of a compound of formula X as defined in h) above which comprises rearrangement of a compound of formula IX as defined in e) above
- j) a process for the production of a compound of formula IX as defined in e) above which comprises alkylation of a compound of formula XII

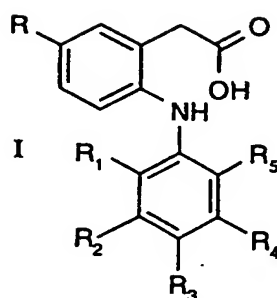


with 2-chloro-N-(4-methylphenyl)acetamide or 2 chloro-N-(4-ethylphenyl)acetamide

- k) a process for the production of a compound of formula VIII as defined in d) above which comprises alkylation of a compound of formula XII as defined in j) above with 2-chloro-N-(4-methylphenyl)acetamide or 2 chloro-N-(4-ethylphenyl)acetamide followed by rearrangement and cleavage
- l) a process for the production of a compound of formula VIII as defined in d) above comprising oxidation of the corresponding compound of formula XIII (or a tautomer thereof)



- m) a process for the production of a compound of formula XIII as defined in l) above which comprises coupling 1-methoxy-4-methylcyclohexa-1,4-diene or 1-methoxy-4-ethylcyclohexa-1,4-diene with an aniline derivative of formula IV as defined in c) above, and
 - n) a process for the production of a compound of formula VIII as defined in d) above comprising coupling 1-methoxy-4-methylcyclohexa-1,4-diene or 1-methoxy-4-ethylcyclohexa-1,4-diene with an aniline derivative of formula IV as defined in c) above, followed by oxidation, wherein the symbols are as defined in claim 1.
3. A process for the preparation of a compound of formula I as defined in claim 1.



which comprises one or more of processes a) to n) as defined in claim 2 and optionally a process according to claim 1.

4. A process according to claim 3 for the preparation of a compound selected from:

5-methyl-2-(2',4'-dichloro-6'-methylanilino)phenylacetic acid;
 5-methyl-2-(2',3',5',6'-tetrafluoroanilino)phenylacetic acid;
 5-methyl-2-(2',3',4',6'-tetrafluoroanilino)phenylacetic acid;
 5-methyl-2-(2',6'-dichloroanilino)phenylacetic acid;
 5-methyl-2-(2',6'-dichloroanilino)phenylacetic acid, potassium salt;
 5-methyl-2-(2',6'-dichloroanilino)phenylacetic acid, sodium salt;
 5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid;
 5-methyl-2-(2',6'-dichloro-4'-methylanilino)phenylacetic acid;
 5-methyl-2-(2'-chloro-6'-methylanilino)phenylacetic acid;
 5-methyl-2-(2',4'-difluoro-6'-chloroanilino)phenylacetic acid;
 5-methyl-2-(2'-fluoro-4',6'-dichloroanilino)phenylacetic acid;
 5-methyl-2-(2'-chloro-4'-fluoro-6'-methylanilino)phenylacetic acid;
 5-ethyl-2-(2'-fluoro-6'-chloroanilino)phenylacetic acid;
 5-ethyl-2-(2'-chloro-6'-methylanilino)phenylacetic acid;
 5-ethyl-2-(2',3',6'-trifluoroanilino)phenylacetic acid;

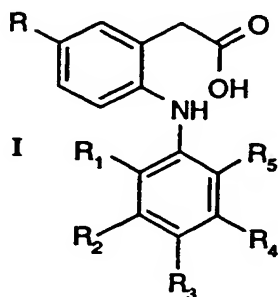
5-ethyl-2-(2',3',5',6'-tetrafluoro-4'-ethoxyanilino)phenylacetic acid;
5-ethyl-2-(2'-chloro-4',6'-difluoroanilino)phenylacetic acid;
5-ethyl-2-(2',4'-dichloro-6'-fluoroanilino)phenylacetic acid;
5-ethyl-2-(2',4'-dichloro-6'-methylanilino)phenylacetic acid;
5-ethyl-2-(2'-fluoro-4'-chloro-6'-methylanilino)phenylacetic acid;
5-ethyl-2-(2',4'-difluoro-6'-methylanilino)phenylacetic acid;
5-ethyl-2-(2'-chloro-4'-fluoro-6'-methylanilino)phenylacetic acid;
5-methyl-2-(2'-chloro-4'-hydroxy-6'-fluoroanilino)phenylacetic acid;
5-methyl-2-(2'-fluoro-6'-trifluoromethylanilino)phenylacetic acid, and
5-methyl-2-(2',4'-dichloro-6'-trifluoromethylanilino)phenylacetic acid,
and pharmaceutically acceptable salts thereof; and pharmaceutically acceptable
prodrug esters thereof.

5. A process according to claim 3 for the preparation of a compound selected from:

5-methyl-2-(2', 3', 4', 6'-tetrafluoroanilino)phenylacetic acid;
5-methyl-2-(2',6'-dichloroanilino)phenylacetic acid;
5-methyl-2-(2'-chloro-6'fluoroanilino)phenylacetic acid;
5-methyl-2-(2',6'-dichloro-4'-methylanilino)phenylacetic acid;
5-methyl-2-(2'-chloro-6'-methylanilino)phenylacetic acid;
5-methyl-2-(2'-chloro-4'-fluoro-6'-methylanilino)phenylacetic acid;
5-ethyl-2-(2'-fluoro-6'-chloroanilino)phenylacetic acid;
5-ethyl-2-(2'-chloro-6'-methylanilino)phenylacetic acid;
5-ethyl-2-(2',3',6'-trifluoroanilino)phenylacetic acid, and
5-ethyl-2-(2',4'-dichloro-6'-methylanilino)phenylacetic acid,

and pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

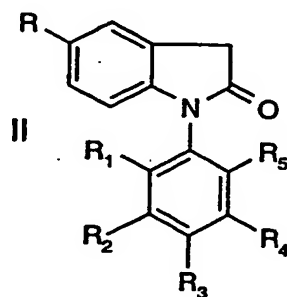
6. A compound of formula I, as defined in claim 1



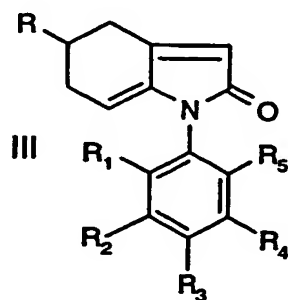
or a pharmaceutically acceptable salt thereof, or a pharmaceutically acceptable prodrug ester thereof, when prepared by a process as defined in claim 3.

7. A compound selected from

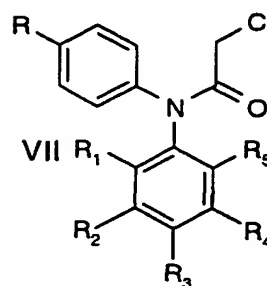
- a) a compound of formula II



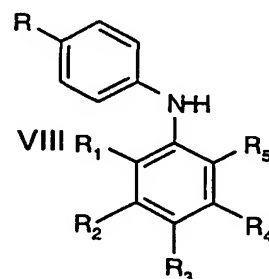
- b) a compound of formula III



- c) a compound of formula VII

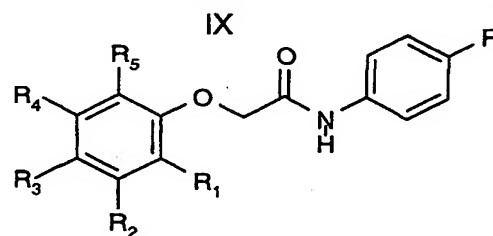


- d) a compound of formula VIII

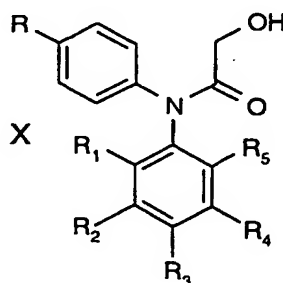


provided that R is not methyl when all of R₁, R₂, R₃, R₄ and R₅ are fluoro;

- e) a compound of formula IX



- f) a compound of formula X



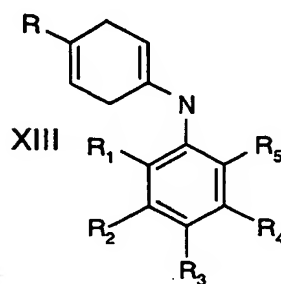
or

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(67)

g) a compound of formula XIII



wherein the symbols are as defined in claim 1.